

**K983897 CERVICAL ANESTHESIA NEEDLE,
CONTRACERVICAL ANESTHESIA NEEDLE, MODEL #'S
720209 30GAU X 4 MM, 720210 30GAU X 6MM, 720208G**

Feb 1, 1999
90 days to decision

K983897 · Product code: **HEE** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k983897/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Anesthesia, Paracervical (HEE)
Date received	Nov 3, 1998
Decision date	Feb 1, 1999
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Ri Mos. S.R.L.
Location	Concord, MA, US
Contact	ADENA S RIEMER
510(k) history	4 submissions · 4 cleared · 1999-1999

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k983897/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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